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NEWS	1		Web Page for STN Seminar Schedule - N. America
NEWS	2	DEC 01	ChemPort single article sales feature unavailable
NEWS	3	JUN 01	CAS REGISTRY Source of Registration (SR) searching enhanced on STN
NEWS	4	JUN 26	NUTRACEUT and PHARMAML no longer updated
NEWS	5	JUN 29	IMSCOPROFILE now reloaded monthly
NEWS	6	JUN 29	EPFULL adds Simultaneous Left and Right Truncation (SLART) to AB, MCLM, and TI fields
NEWS	7	JUL 09	PATDPAFULL adds Simultaneous Left and Right Truncation (SLART) to AB, CLM, MCLM, and TI fields
NEWS	8	JUL 14	USGENE enhances coverage of patent sequence location (PSL) data
NEWS	9	JUL 27	CA/CAplus enhanced with new citing references
NEWS	10	JUL 16	GBFULL adds patent backfile data to 1855
NEWS	11	JUL 21	USGENE adds bibliographic and sequence information
NEWS	12	JUL 28	EPFULL adds first-page images and applicant-cited references
NEWS	13	JUL 28	INPADOCDB and INPAFAMDB add Russian legal status data
NEWS	14	AUG 08	Improve STN by completing a survey and be entered to win a gift card
NEWS	15	AUG 10	Time limit for inactive STN sessions doubles to 40 minutes
NEWS	16	AUG 17	CAS REGISTRY, the Global Standard for Chemical Research, Approaches 50 Millionth Registration Milestone
NEWS	17	AUG 18	COMPENDEX indexing changed for the Corporate Source (CS) field

NEWS EXPRESS MAY 26 09 CURRENT WINDOWS VERSION IS V8.4,
AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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* See NEWS 14 for details or go directly to the survey at:            *
* http://www.zoomerang.com/Survey/?p=WEB229H4S8Q5UL                   *
*                                                                     *
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FILE 'HOME' ENTERED AT 12:52:36 ON 18 AUG 2009

=> file  reg
COST IN U.S. DOLLARS                                  SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                       0.44       0.44

FILE 'REGISTRY' ENTERED AT 12:53:40 ON 18 AUG 2009
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STRUCTURE FILE UPDATES:   17 AUG 2009  HIGHEST RN 1174495-28-3
DICTIONARY FILE UPDATES:  17 AUG 2009  HIGHEST RN 1174495-28-3

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TSCA INFORMATION NOW CURRENT THROUGH June 26, 2009.

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Uploading C:\Program Files\STNEXP\Queries\10591722b.str
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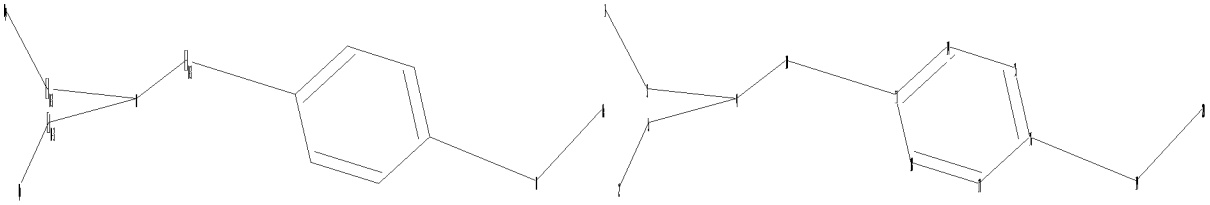

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chain nodes :
1  2  3  4  5  10  19  20
ring nodes :
11  14  15  16  17  18
chain bonds :
1-3  2-5  3-4  4-5  4-10  10-11  16-19  19-20
ring bonds :
11-14  11-18  14-15  15-16  16-17  17-18
```

exact/norm bonds :
1-3 2-5 3-4 4-5 4-10 16-19 19-20
exact bonds :
10-11
normalized bonds :
11-14 11-18 14-15 15-16 16-17 17-18

Match level :
1:Atom 2:Atom 3:CLASS 4:CLASS 5:CLASS 10:CLASS 11:Atom 14:Atom 15:Atom
16:Atom 17:Atom 18:Atom 19:CLASS 20:CLASS

L1   STRUCTURE UPLOADED

=> s l1 sss full
FULL SEARCH INITIATED 12:56:15 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 3619493 TO ITERATE

 43.0% PROCESSED 1556648 ITERATIONS   394 ANSWERS

 50.6% PROCESSED 1829879 ITERATIONS   395 ANSWERS

 55.0% PROCESSED 1992259 ITERATIONS   396 ANSWERS

 55.3% PROCESSED 2000000 ITERATIONS   396 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.53

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**
                        BATCH   **INCOMPLETE**
PROJECTED ITERATIONS: 3619493 TO 3619493
PROJECTED ANSWERS: 636 TO   796

L2   396 SEA SSS FUL L1

=> file capl
COST IN U.S. DOLLARS   SINCE FILE   TOTAL
                                                         ENTRY   SESSION
FULL ESTIMATED COST   188.28   188.72

FILE 'CAPLUS' ENTERED AT 12:57:31 ON 18 AUG 2009
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FILE COVERS 1907 - 18 Aug 2009 VOL 151 ISS 8
FILE LAST UPDATED: 17 Aug 2009 (20090817/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2009

USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2009

CAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2009.

CAS Information Use Policies apply and are available at:

http://www.cas.org/legal/infopolicy.html

This file contains CAS Registry Numbers for easy and accurate substance identification.

The ALL, BIB, MAX, and STD display formats in the CA/CAplus family of databases have been updated to include new citing references information. This enhancement may impact record import into database management software. For additional information, refer to NEWS 9.

```
=> s l2
L3          16 L2

=> d l3 1-16 ibib hitstr

L3   ANSWER 1 OF 16  CAPLUS  COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER:        2009:875995  CAPLUS
DOCUMENT NUMBER:         151:115083
TITLE:                   Method using lifespan-altering compounds for altering
                         the lifespan of eukaryotic organisms, and screening
                         for such compounds
INVENTOR(S):             Goldfarb, David Scott
PATENT ASSIGNEE(S):      University of Rochester, USA
SOURCE:                  U.S. Pat. Appl. Publ., 57pp.
                         CODEN: USXXCO
DOCUMENT TYPE:           Patent
LANGUAGE:                English
PATENT INFORMATION:

     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
     ---------------     ----   --------    --------------------   --------
     US 20090163545 A1          20090625   US 2008-XQ341615       20081222
PRIORITY APPLN. INFO.:                     US 2007-16362P         20071221
                                           US 2008-23801P         20080125
IT   1032762-38-1
     RL: PAC (Pharmacological activity); BIOL (Biological study)
        (method using lifespan-altering compds. for altering lifespan of
        eukaryotic organisms, and screening for such compds.)
RN   1032762-38-1  CAPLUS
CN   1H-Indole-3-acetamide, N-[4-(acetylamino)phenyl]-N-[2-(cyclohexylamino)-1-
     (2-furanyl)-2-oxoethyl]-  (CA INDEX NAME)
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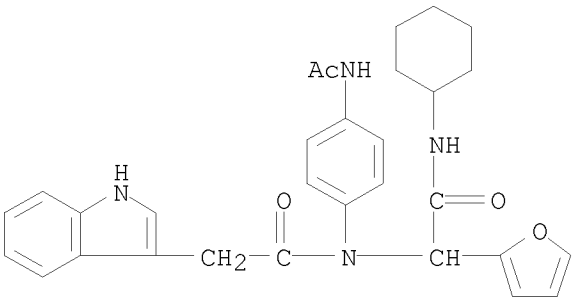
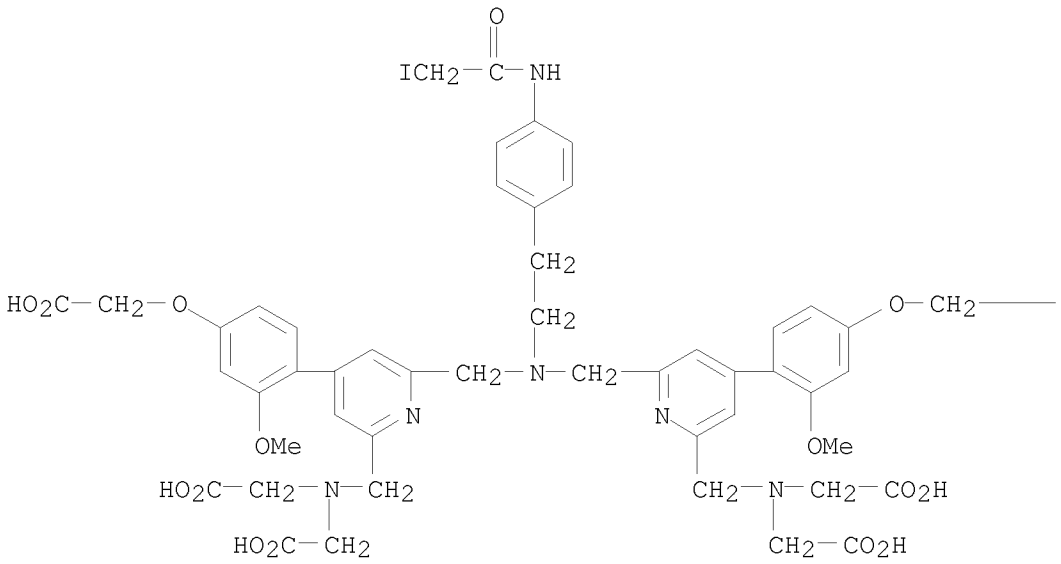


L3 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2009:357199 CAPLUS
DOCUMENT NUMBER: 150:364629
TITLE: Lanthanide pyridine iminodicarboxylate chelate complexes as fluorescent markers for peptides and oligonucleotides
PATENT ASSIGNEE(S): Wallac Oy, Finland
SOURCE: Ger. Gebrauchsmusterschrift, 7pp.
CODEN: GGXXFR
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 202008013315	U1	20090326	DE 2008-202008013315	20081007
PRIORITY APPLN. INFO.:			FI 2007-493U U	20071217
OTHER SOURCE(S):	MARPAT 150:364629			

IT 1133438-12-6P 1133438-13-7P
RL: PEP (Physical, engineering or chemical process); PRPH (Prophetic); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); PROC (Process); RACT (Reactant or reagent)
(preparation of lanthanide pyridine iminodicarboxylate chelate complexes as fluorescent markers for peptides and oligonucleotides)
RN 1133438-12-6 CAPLUS
CN INDEX NAME NOT YET ASSIGNED

PAGE 1-A

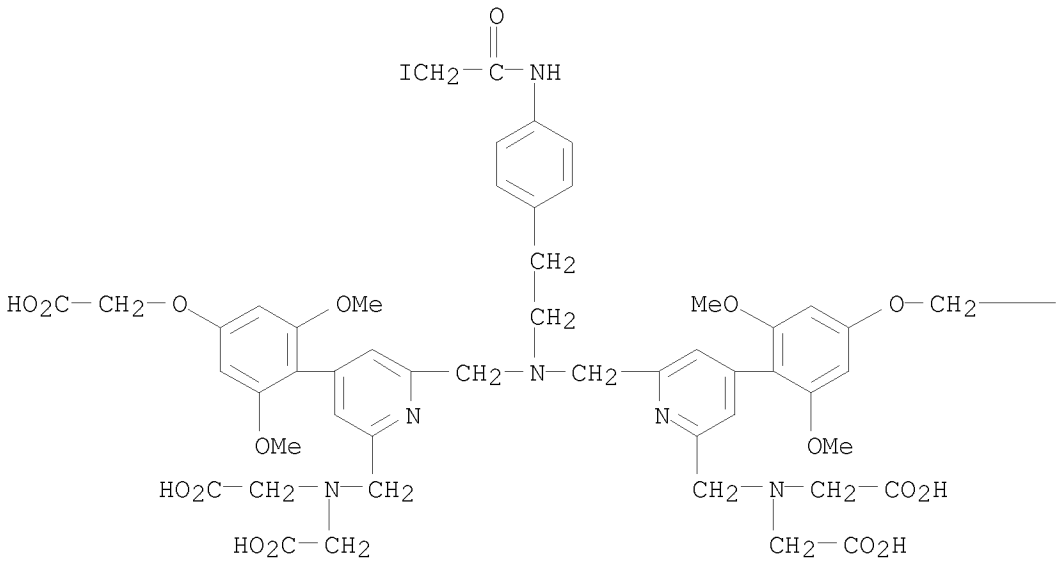


PAGE 1-B

— CO_2H

RN 1133438-13-7 CAPLUS
CN INDEX NAME NOT YET ASSIGNED

PAGE 1-A



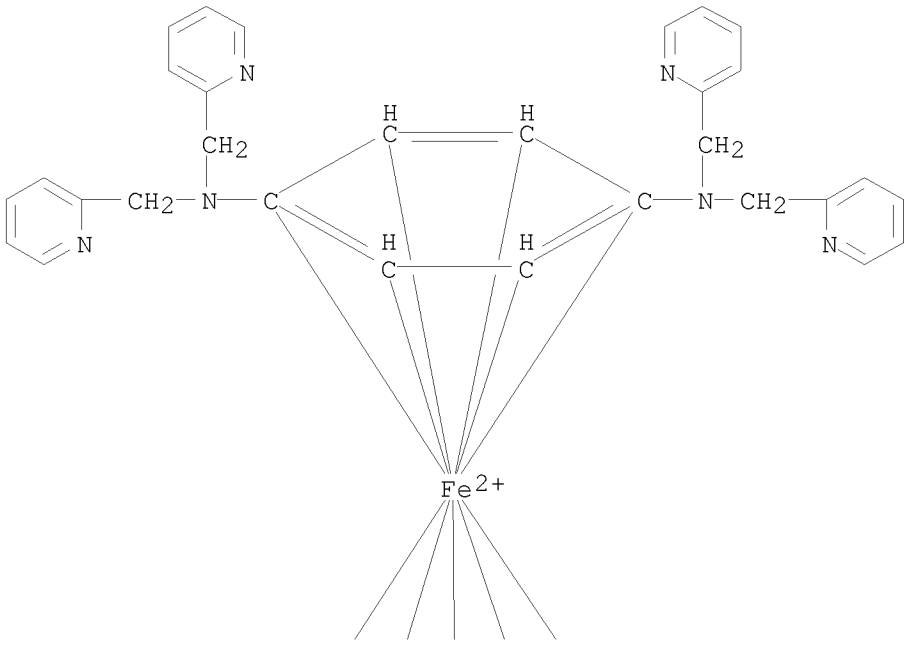
PAGE 1-B

— CO_2H

L3 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2009:247359 CAPLUS
DOCUMENT NUMBER: 150:422650
TITLE: Ultrasound-promoted aromatic nucleophilic substitution of dichlorobenzene iron(II) complexes
AUTHOR(S): Raouafi, Noureddine; Belhadj, Nadra; Boujlel, Khaled; Ourari, Ali; Amatore, Christian; Maisonhaute, Emmanuel; Schoellhorn, Bernd
CORPORATE SOURCE: Departement de Chimie, Faculte des Sciences de Tunis, Universite de Tunis El Manar, Tunis, 2092, Tunisia
SOURCE: Tetrahedron Letters (2009), 50(15), 1720-1722
CODEN: TELEAY; ISSN: 0040-4039
PUBLISHER: Elsevier Ltd.
DOCUMENT TYPE: Journal
LANGUAGE: English
IT 1142881-34-2P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of anilines by ultrasound-promoted aromatic nucleophilic substitution of chlorobenzene iron complexes)

RN 1142881-34-2 CAPLUS
CN Iron(1+), (η5-2,4-cyclopentadien-1-yl)[(1,2,3,4,5,6-η)-N1,N1,N4,N4-tetrakis(2-pyridinylmethyl)-1,4-benzenediamine]- (CA INDEX NAME)

PAGE 1-A



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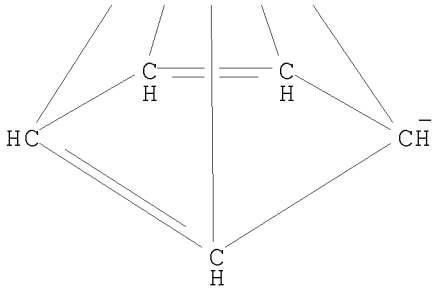


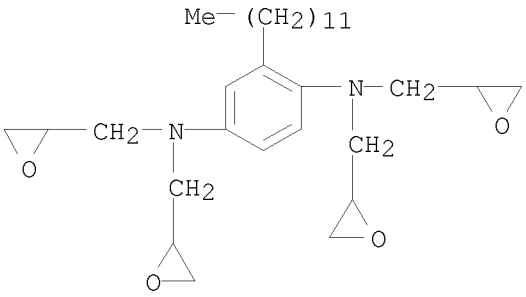
REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:820982 CAPLUS
DOCUMENT NUMBER: 149:211889
TITLE: Vertical-alignment liquid crystal aligning agents and vertical-alignment mode liquid crystal display elements
INVENTOR(S): Kumagaya, Tsutomu; Nishikawa, Michinori
PATENT ASSIGNEE(S): Jsr Corporation, Japan
SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 34pp.
CODEN: CNXXEV
DOCUMENT TYPE: Patent
LANGUAGE: Chinese

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 101210184	A	20080702	CN 2007-10305939	20071228
JP 2008181102	A	20080807	JP 2007-323414	20071214
KR 2008063148	A	20080703	KR 2007-138689	20071227
PRIORITY APPLN. INFO.:			JP 2006-354460 A	20061228

IT 1041184-76-2
RL: TEM (Technical or engineered material use); USES (Uses)
(liquid crystal aligning agents for vertical-alignment mode liquid crystal displays)
RN 1041184-76-2 CAPLUS
CN 1,4-Benzenediamine, 2-dodecyl-N1,N1,N4,N4-tetrakis(2-oxiranylmethyl)- (CA INDEX NAME)



L3 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:736395 CAPLUS
DOCUMENT NUMBER: 149:79490
TITLE: Carboxamides as ion channel modulators and their preparation, pharmaceutical compositions and use in the treatment of diseases
INVENTOR(S): Galullo, Vincent; Zelle, Robert; Mazdiyasni, Hormoz; Baker, Christopher Todd; Will, Paul; Guo, Jinsong; Fensome, Andrew; Soenen, Danielle; Kern, Jeffrey Curtis; Moore, William Jay; Melenski, Edward George; Kaplan, Justin; Sabatucci, Joseph Peter
PATENT ASSIGNEE(S): Wyeth, John, and Brother Ltd., USA
SOURCE: PCT Int. Appl., 312pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008073461	A2	20080619	WO 2007-US25416	20071211
WO 2008073461	A3	20080912		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,

IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

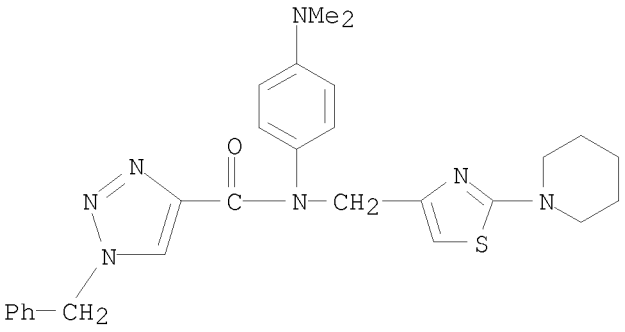
PRIORITY APPLN. INFO.: US 2006-874133P P 20061211
US 2006-874152P P 20061211
US 2006-874179P P 20061211

OTHER SOURCE(S): MARPAT 149:79490

IT 1033831-47-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of carboxamide compds. as ion channel modulators useful in treatment of diseases)

RN 1033831-47-8 CAPLUS

CN 1H-1,2,3-Triazole-4-carboxamide, N-[4-(dimethylamino)phenyl]-1-(phenylmethyl)-N-[[2-(1-piperidinyl)-4-thiazolyl]methyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS)

L3 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:733525 CAPLUS

DOCUMENT NUMBER: 149:53863

TITLE: Preparation of of N,N-substituted 3-aminopyrrolidine compounds useful as monoamines reuptake inhibitors

INVENTOR(S): Kurimura, Muneaki; Taira, Shinichi; Tomoyasu, Takahiro; Ito, Nobuaki; Tai, Kuninori; Takemura, Noriaki; Matsuzaki, Takayuki; Menjo, Yasuhiro; Miyamura, Shin; Sakurai, Yoji; Watabe, Akihito; Sakata, Yasuyo; Masumoto, Takumi; Akazawa, Kohei; Sugino, Haruhiko; Amada, Naoki; Ohashi, Satoshi; Shinohara, Tomokazu; Sasaki, Hirofumi; Morita, Chisako; Yamashita, Junko; Nakajima, Satoko

PATENT ASSIGNEE(S): Ohtsuka Pharmaceutical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 221pp.
CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2008137997	A	20080619	JP 2007-292386	20071109

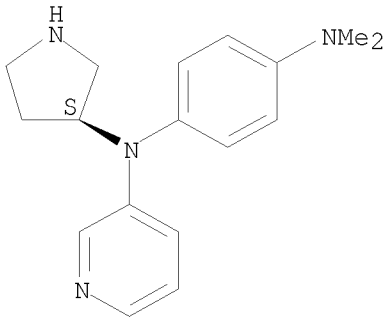
PRIORITY APPLN. INFO.: JP 2006-305573 A 20061110

OTHER SOURCE(S): MARPAT 149:53863

IT 914997-33-4P 914997-67-4P 914997-70-9P
915000-91-8P 915001-14-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of aminopyrrolidine compds. as monoamines reuptake inhibitors with sufficient therapeutic effects after short-term administration)

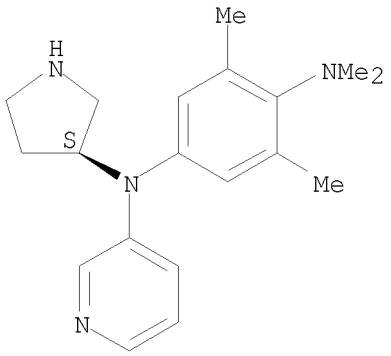
RN 914997-33-4 CAPLUS
CN 1,4-Benzenediamine, N1,N1-dimethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



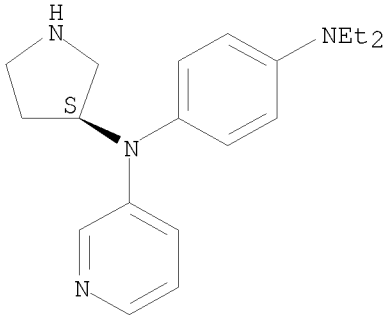
RN 914997-67-4 CAPLUS
CN 1,4-Benzenediamine, N1,N1,2,6-tetramethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



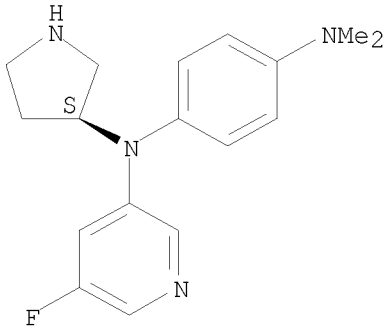
RN 914997-70-9 CAPLUS
CN 1,4-Benzenediamine, N1,N1-diethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



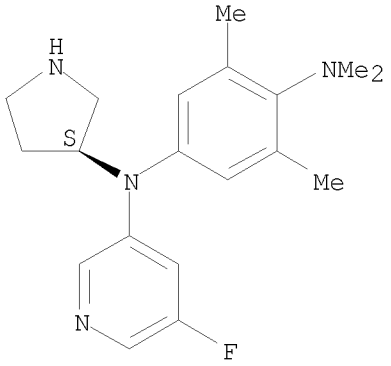
RN 915000-91-8 CAPLUS
CN 1,4-Benzenediamine, N1-(5-fluoro-3-pyridinyl)-N4,N4-dimethyl-N1-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



RN 915001-14-8 CAPLUS
CN 1,4-Benzenediamine, N4-(5-fluoro-3-pyridinyl)-N1,N1,2,6-tetramethyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.

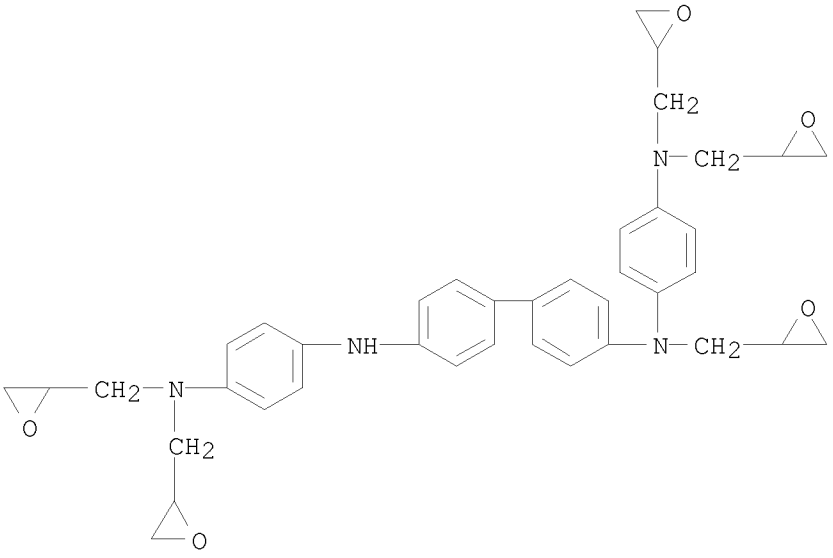


L3 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:447206 CAPLUS
DOCUMENT NUMBER: 148:506743
TITLE: Liquid crystal alignment agent and liquid crystal display element
INVENTOR(S): Yasuda, Hiroyuki; Hayashi, Eiji; Nishikawa, Michinori

PATENT ASSIGNEE(S): Jsr Corporation, Japan
SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 43pp.
CODEN: CNXXEV
DOCUMENT TYPE: Patent
LANGUAGE: Chinese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 101153995	A	20080402	CN 2007-10151310	20070924
KR 2008028320	A	20080331	KR 2007-96550	20070921
JP 2008107811	A	20080508	JP 2007-247604	20070925
PRIORITY APPLN. INFO.:			JP 2006-261196 A	20060926

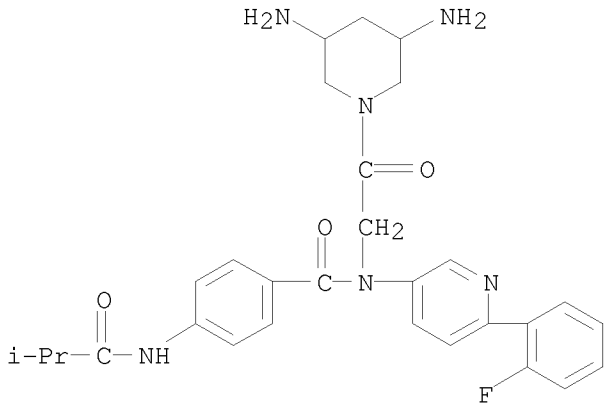
IT 1020839-25-1P
RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
(liquid crystal alignment agent and liquid crystal display element)
RN 1020839-25-1 CAPLUS
CN [1,1'-Biphenyl]-4,4'-diamine, N4,N4'-bis[4-[bis(2-oxiranylmethyl)amino]phenyl]-N4-(2-oxiranylmethyl)- (CA INDEX NAME)



L3 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:124359 CAPLUS
DOCUMENT NUMBER: 148:191838
TITLE: Preparation of substituted aniline derivatives as antifungal agents
INVENTOR(S): Carr, Andrew David; Neuss, Judi Charlotte; Orchard, Michael Glen; Porter, David William
PATENT ASSIGNEE(S): Ucb Pharma S.A., Belg.
SOURCE: PCT Int. Appl., 111pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

WO 2008012524 A1 20080131 WO 2007-GB2815 20070724
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
AU 2007279092 A1 20080131 AU 2007-279092 20070724
CA 2658913 A1 20080131 CA 2007-2658913 20070724
US 20080045497 A1 20080221 US 2007-782337 20070724
EP 2046739 A1 20090415 EP 2007-766352 20070724
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS
PRIORITY APPLN. INFO.: GB 2006-14677 A 20060724
GB 2006-14678 A 20060724
GB 2007-4645 A 20070309
GB 2007-4648 A 20070309
WO 2007-GB2815 W 20070724
OTHER SOURCE(S): MARPAT 148:191838
IT 1013324-35-0P
RL: PAC (Pharmacological activity); PRPH (Prophetic); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of substituted aniline derivs. as antifungal agents)
RN 1013324-35-0 CAPLUS
CN Benzamide, N-[2-(3,5-diamino-1-piperidinyl)-2-oxoethyl]-N-[6-(2-fluorophenyl)-3-pyridinyl]-4-[(2-methyl-1-oxopropyl)amino]- (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2007:1364437 CAPLUS
DOCUMENT NUMBER: 148:33637
TITLE: Substituted quinolones as ATP-utilizing enzyme inhibitors and their preparation, compositions, and uses thereof

INVENTOR(S): Dickson, John K.; Chen, Ke; Hodge, Carl Nicholas
PATENT ASSIGNEE(S): Amphora Discovery Corporation, USA
SOURCE: PCT Int. Appl., 143pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007136592	A2	20071129	WO 2007-US11484	20070510
WO 2007136592	A3	20080228		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
CA 2652634	A1	20071129	CA 2007-2652634	20070510
US 20070287706	A1	20071213	US 2007-803140	20070510
EP 2040711	A2	20090401	EP 2007-794818	20070510
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS				

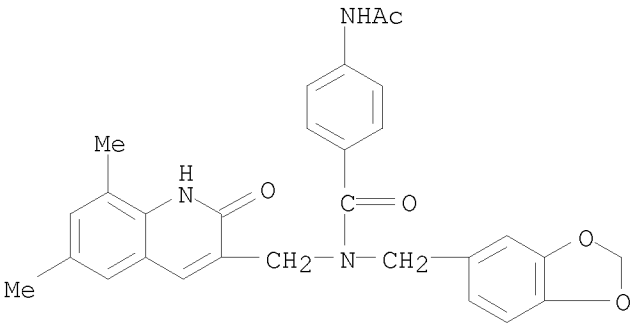
PRIORITY APPLN. INFO.: US 2006-801881P P 20060518
WO 2007-US11484 W 20070510

OTHER SOURCE(S): MARPAT 148:33637

IT 958454-68-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of substituted quinolones as ATP-utilizing enzyme inhibitors useful in the treatment of diseases)

RN 958454-68-7 CAPLUS

CN Benzamide, 4-(acetylamino)-N-(1,3-benzodioxol-5-ylmethyl)-N-[(1,2-dihydro-6,8-dimethyl-2-oxo-3-quinolinyl)methyl]- (CA INDEX NAME)

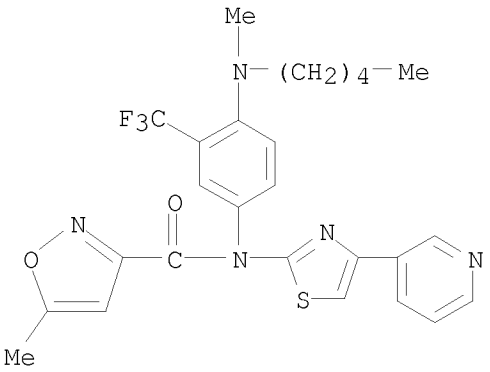


OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L3 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2007:1029651 CAPLUS
DOCUMENT NUMBER: 147:365486
TITLE: Preparation of 2-(phenylamino)thiazole derivatives as inhibitors of viral replication for the treatment of hepatitis C infection
INVENTOR(S): Zhang, Suoming; Phadke, Avinash; Wang, Xiangzhu; Liu, Cuixian
PATENT ASSIGNEE(S): Achillion Pharmaceuticals, Inc., USA
SOURCE: PCT Int. Appl., 134pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007103550	A2	20070913	WO 2007-US6023	20070308
WO 2007103550	A3	20071108		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
AU 2007223797	A1	20070913	AU 2007-223797	20070308
CA 2645072	A1	20070913	CA 2007-2645072	20070308
US 20070213301	A1	20070913	US 2007-683749	20070308
EP 1996565	A2	20081203	EP 2007-752705	20070308
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS				
JP 2009529059	T	20090813	JP 2008-558416	20070308

PRIORITY APPLN. INFO.: US 2006-780609P P 20060308
WO 2007-US6023 W 20070308
OTHER SOURCE(S): CASREACT 147:365486; MARPAT 147:365486
IT 949117-20-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of (phenylamino)thiazoles as inhibitors of viral replication for treatment of hepatitis C infection)
RN 949117-20-8 CAPLUS
CN 3-Isoxazolecarboxamide, 5-methyl-N-[4-(methylpentylamino)-3-(trifluoromethyl)phenyl]-N-[4-(3-pyridinyl)-2-thiazolyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L3 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2007:1028755 CAPLUS
DOCUMENT NUMBER: 147:365493
TITLE: Heterobicyclic pyrazole compounds as Met tyrosine kinase inhibitors and their preparation and use
INVENTOR(S): Blake, James F.; Boyd, Steven Armen; Cohen, Frederick; De Meese, Jason; Fong, Kin Chiu; Gaudino, John J.; Kaplan, Tomas; Marlow, Allison L.; Seo, Jeongbeob; Thomas, Allen A.; Tian, Hongqi; Young, Wendy B.
PATENT ASSIGNEE(S): Array Biopharma Inc., USA; Genentech, Inc.
SOURCE: PCT Int. Appl., 273 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007103308	A2	20070913	WO 2007-US5583	20070306
WO 2007103308	A3	20080207		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
AU 2007224020	A1	20070913	AU 2007-224020	20070306
CA 2645137	A1	20070913	CA 2007-2645137	20070306
US 20070238726	A1	20071011	US 2007-714342	20070306
EP 2001880	A2	20081217	EP 2007-752297	20070306
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS				
JP 2009529047	T	20090813	JP 2008-558335	20070306
MX 2008011220	A	20080911	MX 2008-11220	20080902
IN 2008KN03882	A	20090227	IN 2008-KN3882	20080924

NO 2008004183	A	20081124	NO 2008-4183		20081006
KR 2008110783	A	20081219	KR 2008-724415		20081006
CN 101437820	A	20090520	CN 2007-80016155		20081104
PRIORITY APPLN. INFO.:			US 2006-779805P	P	20060307
			US 2006-874832P	P	20061214
			WO 2007-US5583	W	20070306

OTHER SOURCE(S): MARPAT 147:365493

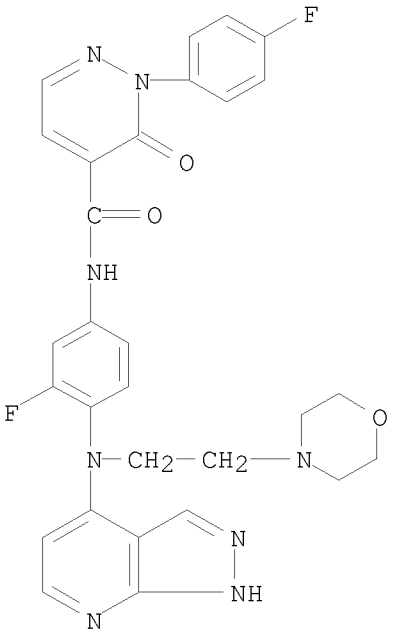
IT 949560-23-0P 949560-28-5P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of heterobicyclic pyrazole compds. as Met tyrosine kinase inhibitors useful in the treatment of diseases)

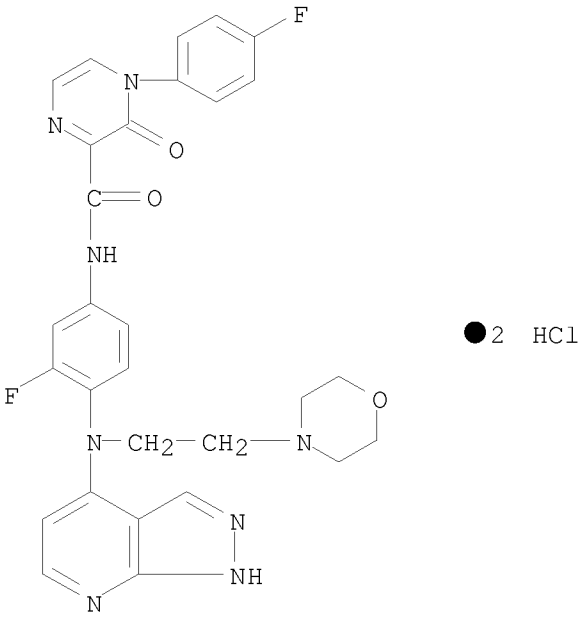
RN 949560-23-0 CAPLUS

CN 4-Pyridazinecarboxamide, N-[3-fluoro-4-[[2-(4-morpholinyl)ethyl]-1H-pyrazolo[3,4-b]pyridin-4-ylamino]phenyl]-2-(4-fluorophenyl)-2,3-dihydro-3-oxo- (CA INDEX NAME)



RN 949560-28-5 CAPLUS

CN 2-Pyrazinecarboxamide, N-[3-fluoro-4-[[2-(4-morpholinyl)ethyl]-1H-pyrazolo[3,4-b]pyridin-4-ylamino]phenyl]-4-(4-fluorophenyl)-3,4-dihydro-3-oxo-, hydrochloride (1:2) (CA INDEX NAME)

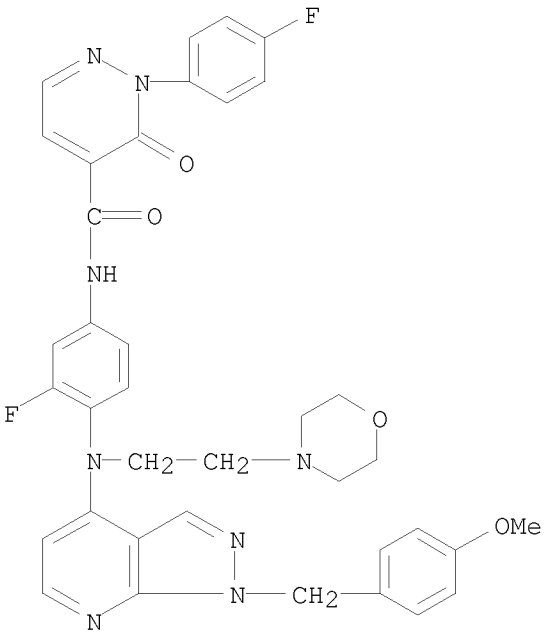


● 2 HCl

IT 949560-27-4P, N-[4-[[1-(4-Methoxybenzyl)-1H-pyrazolo[3,4-b]pyridin-4-yl](2-morpholinoethyl)amino]-3-fluorophenyl]-2-(4-fluorophenyl)-3-oxo-2,3-dihydropyridazine-4-carboxamide
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of heterobicyclic pyrazole compds. as Met tyrosine kinase inhibitors useful in the treatment of diseases)

RN 949560-27-4 CAPLUS

CN 4-Pyridazinecarboxamide, N-[3-fluoro-4-[[1-[(4-methoxyphenyl)methyl]-1H-pyrazolo[3,4-b]pyridin-4-yl][2-(4-morpholinyl)ethyl]amino]phenyl]-2-(4-fluorophenyl)-2,3-dihydro-3-oxo- (CA INDEX NAME)



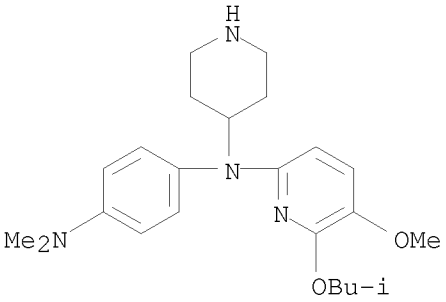
OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L3 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2006:1338137 CAPLUS
DOCUMENT NUMBER: 146:81773
TITLE: Preparation of N-substituted diarylamine analogs as phosphodiesterase 4 inhibitors
INVENTOR(S): Talamas, Francisco Xavier; Caroon, Joan Marie; Dunn, Robert; Hopper, Allen; Kuester, Eric; Schumacher, Richard; Tehim, Ashok
PATENT ASSIGNEE(S): Memory Pharmaceuticals Corporation, USA; F. Hoffmann-La Roche A.-G.
SOURCE: PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006135828	A2	20061221	WO 2006-US22655	20060609
WO 2006135828	A3	20070426		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
AU 2006257863	A1	20061221	AU 2006-257863	20060609
CA 2611562	A1	20061221	CA 2006-2611562	20060609
US 20070049611	A1	20070301	US 2006-449868	20060609
EP 1888528	A2	20080220	EP 2006-784743	20060609
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU				
JP 2008543781	T	20081204	JP 2008-515998	20060609
US 20090118270	A1	20090507	US 2008-329214	20081205

PRIORITY APPLN. INFO.:	US 2005-689060P	P	20050610
	US 2006-449868	A1	20060609
	WO 2006-US22655	W	20060609

OTHER SOURCE(S): MARPAT 146:81773
IT 917098-71-6P, N-(6-Isobutoxy-5-methoxypyridin-2-yl)-N',N'-dimethyl-N-(piperidin-4-yl)benzene-1,4-diamine
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of N-substituted diarylamine analogs as phosphodiesterase 4 inhibitors for treating cognition disorders, inflammation, and other disorders)
RN 917098-71-6 CAPLUS
CN 1,4-Benzenediamine, N1-[5-methoxy-6-(2-methylpropoxy)-2-pyridinyl]-N4,N4-dimethyl-N1-4-piperidinyl- (CA INDEX NAME)



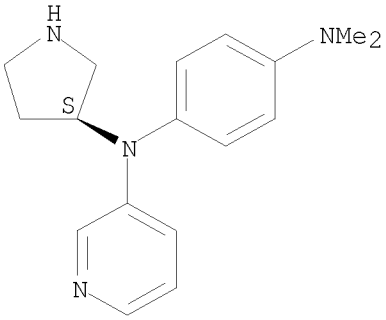
OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)
REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2006:1206758 CAPLUS
DOCUMENT NUMBER: 145:505325
TITLE: Preparation of of N,N-substituted 3-aminopyrrolidine compounds useful as monoamines reuptake inhibitors
INVENTOR(S): Kurimura, Muneaki; Taira, Shinichi; Tomoyasu, Takahiro; Ito, Nobuaki; Tai, Kuninori; Takemura, Noriaki; Matsuzaki, Takayuki; Menjo, Yasuhiro; Miyamura, Shin; Sakurai, Yohji; Watanabe, Akihito; Sakata, Yasuyo; Masumoto, Takumi; Akazawa, Kohei; Sugino, Haruhiko; Amada, Naoki; Ohashi, Satoshi; Shinohara, Tomoichi; Sasaki, Hirofumi; Morita, Chisako; Yamashita, Junko; Nakajima, Satoko
PATENT ASSIGNEE(S): Otsuka Pharmaceutical Co., Ltd., Japan
SOURCE: PCT Int. Appl., 260pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006121218	A1	20061116	WO 2006-JP309988	20060512
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006244851	A1	20061116	AU 2006-244851	20060512
CA 2608184	A1	20061116	CA 2006-2608184	20060512
EP 1881975	A1	20080130	EP 2006-756356	20060512
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
JP 2008540329	T	20081120	JP 2007-552427	20060512
IN 2007DN08276	A	20071123	IN 2007-DN8276	20071026
CN 101175748	A	20080507	CN 2006-80016402	20071112

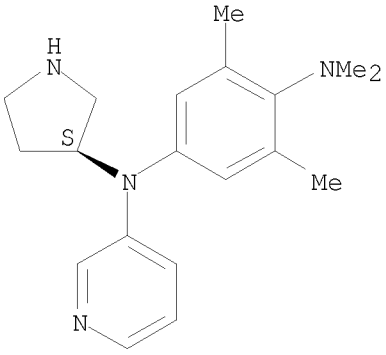
MX 2007014252 A 20080122 MX 2007-14252 20071113
KR 2008008423 A 20080123 KR 2007-729022 20071212
US 20090088406 A1 20090402 US 2008-914183 20080929
PRIORITY APPLN. INFO.: JP 2005-141230 A 20050513
WO 2006-JP309988 W 20060512
OTHER SOURCE(S): MARPAT 145:505325
IT 914997-33-4P 914997-67-4P 914997-70-9P
915000-91-8P 915001-14-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of aminopyrrolidine compds. as monoamines reuptake inhibitors with sufficient therapeutic effects after short-term administration)
RN 914997-33-4 CAPLUS
CN 1,4-Benzenediamine, N1,N1-dimethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



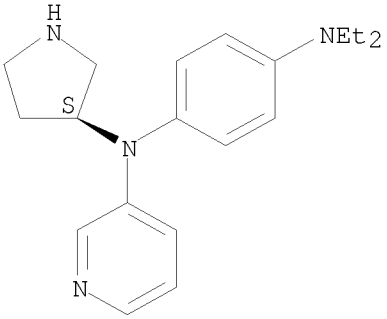
RN 914997-67-4 CAPLUS
CN 1,4-Benzenediamine, N1,N1,2,6-tetramethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



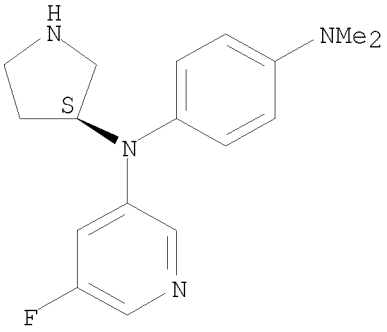
RN 914997-70-9 CAPLUS
CN 1,4-Benzenediamine, N1,N1-diethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



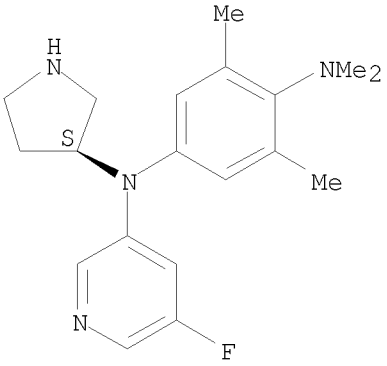
RN 915000-91-8 CAPLUS
CN 1,4-Benzenediamine, N1-(5-fluoro-3-pyridinyl)-N4,N4-dimethyl-N1-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



RN 915001-14-8 CAPLUS
CN 1,4-Benzenediamine, N4-(5-fluoro-3-pyridinyl)-N1,N1,2,6-tetramethyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

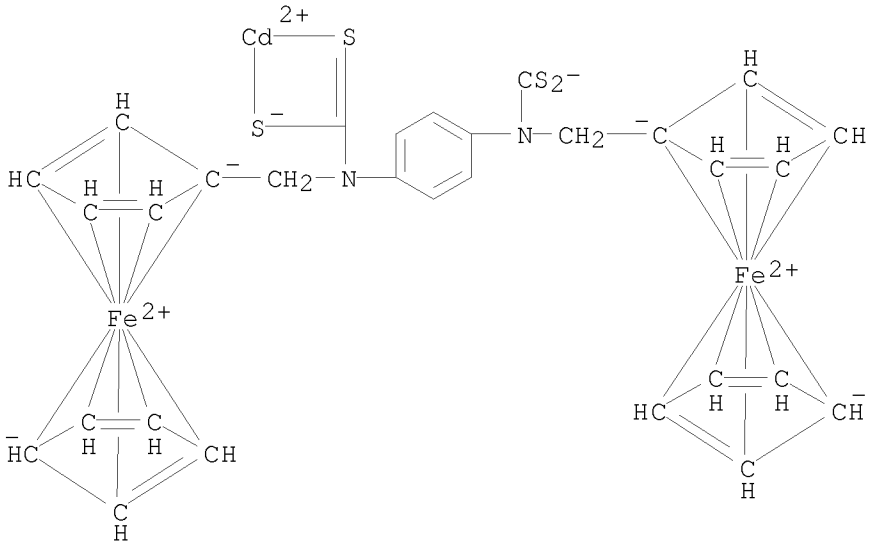
Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2006:109700 CAPLUS
DOCUMENT NUMBER: 145:431181
TITLE: Synthesis and thermal decomposition of cadmium

dithiocarbamate complexes
AUTHOR(S): Thammakan, Nirawan; Somsook, Ekasith
CORPORATE SOURCE: Department of Chemistry, Faculty of Science, Mahidol University, Bangkok, 10400, Thailand
SOURCE: Materials Letters (2006), 60(9-10), 1161-1165
CODEN: MLETDJ; ISSN: 0167-577X
PUBLISHER: Elsevier B.V.
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 145:431181
IT 911824-19-6P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and thermal decomposition of cadmium(II) benzyl- and ferrocenylmethyl-substituted benzenedithiocarbamate polymeric complexes)
RN 911824-19-6 CAPLUS
CN Cadmium, [[[(dithiocarboxy-κS,κS')[4-[(dithiocarboxy)(ferrocenylmethyl)amino]phenyl]amino]methyl]ferrocenato(2-)]- (9CI) (CA INDEX NAME)



OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS)
REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2005:1075811 CAPLUS
DOCUMENT NUMBER: 143:367523
TITLE: Preparation of monosaccharide derivatives as anti-inflammatory agents
INVENTOR(S): Sattigeri, Viswajanani Jitendra; Arora, Sudershan K.; Salman, Mohammad; Palle, Venkata P.; Yadav, Gyan Chand; Tanwar, Madan Pal; Mukherjee, Ashis; Narayanan, Ramamurthy; Rauf, Abdul Rehaman Abdul; Naik, Keshav Prabhakar; Soni, Ajay; Ray, Abhijit; Shirumalla, Raj Kumar; Mookhtiar, Kasim Abbas
PATENT ASSIGNEE(S): Ranbaxy Laboratories Limited, India
SOURCE: PCT Int. Appl., 185 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

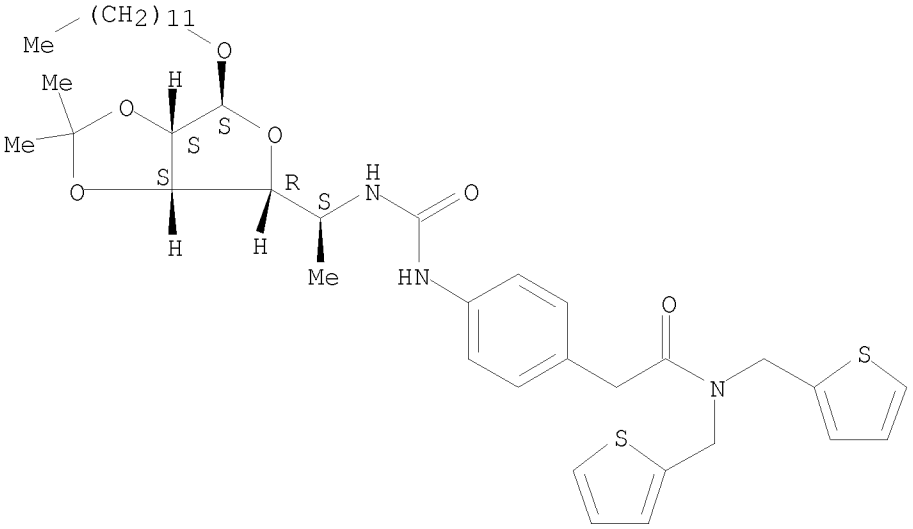
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005092907	A2	20051006	WO 2005-IB803	20050329
WO 2005092907	A3	20060427		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2004-556936P P 20040326
OTHER SOURCE(S): CASREACT 143:367523; MARPAT 143:367523
IT 1043943-91-4
RL: PRPH (Prophetic)
(Preparation of monosaccharide derivatives as anti-inflammatory agents)
RN 1043943-91-4 CAPLUS
CN INDEX NAME NOT YET ASSIGNED

Relative stereochemistry.



L3 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2002:487497 CAPLUS
DOCUMENT NUMBER: 137:78952
TITLE: Preparation of substituted imidazoles, pyrazoles and amides as high affinity C5a receptor modulators
INVENTOR(S): Thurkauf, Andrew; Zhang, Xiaoyan; He, Xia-Shu; Zhao, He; Peterson, John; Maynard, George; Ohliger, Robert
PATENT ASSIGNEE(S): Neurogen Corporation, USA
SOURCE: PCT Int. Appl., 609 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002049993	A2	20020627	WO 2000-US26816	20000929
WO 2002049993	A3	20030220		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2420215	A1	20020627	CA 2000-2420215	20000929
AU 2000076225	A	20020701	AU 2000-76225	20000929
EP 1322309	A2	20030702	EP 2000-965522	20000929
EP 1322309	B1	20080813		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				
ZA 2003001160	A	20040212	ZA 2003-1160	20000929
BR 2000017338	A	20040427	BR 2000-17338	20000929
JP 2004525873	T	20040826	JP 2002-551496	20000929
AU 2000276225	B2	20080710	AU 2000-276225	20000929
AT 404553	T	20080815	AT 2000-965522	20000929
NO 2003001370	A	20030530	NO 2003-1370	20030326
MX 2003002788	A	20041213	MX 2003-2788	20030328

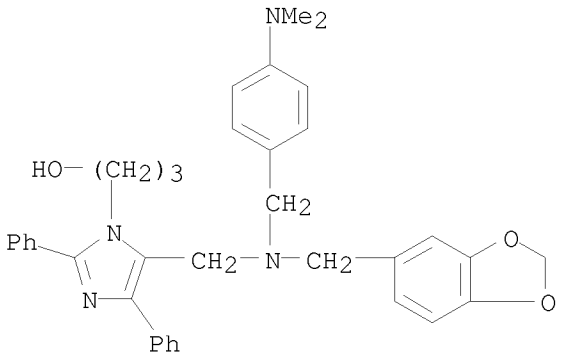
PRIORITY APPLN. INFO.:	US 2000-227454P	P	20000823
	US 1999-156390P	P	19990928
	US 2000-202749P	P	20000508
	US 2000-212449P	P	20000616
	US 2000-221787P	P	20000731
	US 2000-224036P	P	20000809
	WO 2000-US26816	W	20000929

OTHER SOURCE(S): MARPAT 137:78952

IT 1106056-27-2
RL: PRPH (Prophetic)
(Preparation of substituted imidazoles, pyrazoles and amides as high affinity C5a receptor modulators)

RN 1106056-27-2 CAPLUS

CN 1H-Imidazole-1-propanol, 5-[[(1,3-benzodioxol-5-ylmethyl)[[4-(dimethylamino)phenyl]methyl]amino]methyl]-2,4-diphenyl- (CA INDEX NAME)



OS.CITING REF COUNT: 9 THERE ARE 9 CAPLUS RECORDS THAT CITE THIS RECORD (10 CITINGS)

=> file reg
COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 66.24 254.96

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STRUCTURE FILE UPDATES: 17 AUG 2009 HIGHEST RN 1174495-28-3
DICTIONARY FILE UPDATES: 17 AUG 2009 HIGHEST RN 1174495-28-3

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http://www.cas.org/support/stngen/stndoc/properties.html

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